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FILE 'MEDLINE, EMBASE' ENTERED AT 16:49:49 ON 08 MAY 2002
          1180 S PESTIVIRUS
Ll
L2
        181028 S VACCINE
L3
          179 S L1 AND L2
L4
        160557 S ATTENUAT?
L5
            23 S L3 AND L4
L6
            21 DUP REM L5 (2 DUPLICATES REMOVED)
    FILE 'MEDLINE, EMBASE, VETU' ENTERED AT 17:06:00 ON 08 MAY 2002
L7
          1207 S L1
L8
        192850 S L2
           195 S L3
L9
        162607 S L4
L10
L11
            26 S L5
L12
            21 S L6
L13
         1774 S NTR OR NONTRANSLATED REGION
L14
            53 S L13 AND L4
             0 S L14 AND L1
L15
            16 S L8 AND L14
L16
             9 DUP REM L16 (7 DUPLICATES REMOVED)
L17
L18
       1542756 S MUTAT? OR ALTER? OR DELET?
L19
           600 S L18 AND L13
L20
            19 S L19 AND L7
L21
            11 DUP REM L20 (8 DUPLICATES REMOVED)
            18 S VIRUS ATTENUATION AND PROBLEM?
L22
L23
            17 DUP REM L22 (1 DUPLICATE REMOVED)
            67 S VIRUS ATTENUATION AND EFFECTIV?
L24
L25
            67 DUP REM L24 (0 DUPLICATES REMOVED)
      1228275 S REVIEW?
L26
            13 S L25 AND L26
L27
            86 S VIRUS ATTENUATION AND SAF?
L28
            86 DUP REM L28 (0 DUPLICATES REMOVED)
L29
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L30

12 S L29 AND L26

L Number	Hits	Search Text	DB	Time stamp
1	1	pestivirus and (NTR with mutat\$4)	USPAT;	2002/05/08 16:38
			US-PGPUB;	
			EPO; JPO;	
			DERWENT;	
	40	All days with the second second	IBM_TDB	0000/05/00 40:45
2	12	pestivirus with vaccine	USPAT;	2002/05/08 16:45
			US-PGPUB; EPO; JPO;	
			DERWENT;	
			IBM_TDB	
4	26862	vaccine	USPAT;	2002/05/08 16:47
			US-PGPUB;	
			EPO; JPO;	
			DERWENT;	
_			IBM_TDB	
3	15	((nontranslat\$4 adj region\$1) or NTR) with mutat\$4	USPAT;	2002/05/08 16:47
			US-PGPUB;	
			EPO; JPO; DERWENT;	
			IBM_TDB	
5	9	(((nontranslat\$4 adj region\$1) or NTR) with mutat\$4 ) and	USPAT;	2002/05/08 16:48
•		vaccine	US-PGPUB;	
			EPO; JPO;	
			DERWENT,	
			IBM_TDB	

L6 ANSWER 9 OF 21 MEDLINE

ACCESSION NUMBER: 1998235871 MEDLINE

DOCUMENT NUMBER: 98235871 PubMed ID: 9576338

TITLE: Detection of cytopathic bovine viral diarrhea virus in the

ovaries of cattle following immunization with a modified

live bovine viral diarrhea virus vaccine.

AUTHOR: Grooms D L; Brock K V; Ward L A

CORPORATE SOURCE: Ohio Agricultural Research and Development Center, Food

Animal Health Research Program, Wooster 44691, USA.

SOURCE: JOURNAL OF VETERINARY DIAGNOSTIC INVESTIGATION, (1998 Apr)

10 (2) 130-4.

Journal code: A2D; 9011490. ISSN: 1040-6387.

PUB. COUNTRY: United States

Journal; Article; (JOURNAL ARTICLE)

LANGUAGE: English

FILE SEGMENT: Priority Journals

ENTRY MONTH: 199806

ENTRY DATE: Entered STN: 19980625

Last Updated on STN: 19980625 Entered Medline: 19980616

Economic loss from infection with bovine viral diarrhea virus (BVDV) is of AB worldwide concern. The unique pathogenesis and antigenic variability of BVDV have made this virus challenging to control. Vaccination programs are a major component of control and prevention strategies. Both killed and modified live vaccines are commercially available. Choice between killed and modified live vaccines is controversial. Of major concern is the safety of modified live vaccines. Little information is available on their tissue tropism and potential for causing pathology, especially with respect to the reproductive system. The objective of this study was to determine if BVDV could be detected in the ovary of cattle following immunization with a modified live BVDV vaccine. In 2 separate trials, 6 heifers and 4 mature cows were immunized with a modified live BVDV vaccine and ovaries were removed between 7 and 30 days postvaccination. Cytopathic BVDV was isolated from ovaries removed on days 8, 10, and 12. BVDV antigen was detected using immunohistochemistry on days 10-30. These findings are significant because replication of virus in the ovary could cause ovarian dysfunction, resulting in reduced fertility.